# **WEST Search History**

Hide Items Restore Clear Cancel

DATE: Friday, January 06, 2006

Hide?	Hide? <u>Set Name</u> <u>Query</u> <u>H</u>		
	DB=PG	SPB,USPT,EPAB; PLUR=YES; OP=ADJ	
	L12	L11 AND L9	84
	L11	L10 AND L3	323
	L10	L2 AND L1	1146
	L9	L8 or L7	29257
	L8	(435/7.1  435/7.23)![CCLS]	13124
	L7	(530/350)![CCLS]	18276
	L6	(530)![CCLS]	0
	L5	(530)![CCLS]	0
	L4	2A2A9	0
	L3	prostate	38232
	L2	tumor\$ or tumuor\$ or cancer\$ or neoplas\$	180591
	L1	(jakobovits or afar or challita\$ or levin or mitchell or hubert).in	34377

**END OF SEARCH HISTORY** 

# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: December 4, 2005, 09:57:24; Search time 187 Seconds

(without alignments)

1184.208 Million cell updates/sec

Title: US-09-771-312-2

Perfect score: 2694

Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2443163 seqs, 439378781 residues

Total number of hits satisfying chosen parameters: 2443163

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: A Geneseq 21:\*

1: geneseqp1980s:\*

2: geneseqp1990s:\*

3: geneseqp2000s:\*

4: geneseqp2001s:\*

5: geneseqp2002s:\*

6: geneseqp2003as:\*

7: geneseqp2003bs:\*

8: geneseqp2004s:\*

9: geneseqp2005s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	<pre>% Query Match</pre>	Length	DB	ID	Description	
1	2694	100.0	504	4	AAU06524	Aau06524 Prost	ate
2	2694	100.0	528	4	AAB92632	Aab92632 Human	pro
3	2694	100.0	528	5	ABB97288	Abb97288 Novel	hum
4	1808	67.1	376	8	ADR99239	Adr99239 Hypotl	heti
5	825.5	30.6	313	4	ABG08002	Abg08002 Novel	hum
6	608.5	22.6	423	4	ABG23408	Abg23408 Novel	hum
7	591.5	22.0	446	5	ABB75706	Abb75706 Human	pho
8	588.5	21.8	482	5	ABP43772	Abp43772 14 cl	one

```
9
    533.5
            19.8
                    453 5 ABB97561
                                                    Abb97561 Novel hum
10
    530.5
            19.7
                    351 4 AAB94662
                                                    Aab94662 Human pro
    530.5
            19.7
11
                    351 5 ABB97470
                                                    Abb97470 Novel hum
12
      338
            12.5
                    275
                        4 AAB92468
                                                    Aab92468 Human pro
13
    320.5
            11.9
                    223 4 AAM15386
                                                    Aam15386 Peptide #
    320.5 11.9
14
                    223 4 ABB34392
                                                    Abb34392 Peptide #
15
    320.5 11.9
                    223 4 AAM27874
                                                    Aam27874 Peptide #
16
   320.5 11.9
                    223 4 ABB29229
                                                    Abb29229 Peptide #
17
    320.5
          11.9
                    223 4 AAM67577
                                                    Aam67577 Human bon
18
    320.5
            11.9
                    223 4
                           AAM55182
                                                    Aam55182 Human bra
19
    320.5
           11.9
                   223 4 ABG49223
                                                    Abg49223 Human liv
20
    320.5
           11.9
                    223 4 AAM03148
                                                    Aam03148 Peptide #
21
    320.5 11.9
                   223 5 ABG37168
                                                    Abg37168 Human pep
22
    320.5
            11.9
                   223 8 ABO59933
                                                    Abo59933 Human gen
23
    236.5
            8.8
                   123 4
                           AAM95677
                                                    Aam95677 Human rep
24
      196
             7.3
                   1038 7
                           ADC03412
                                                    Adc03412 Rice flow
25
      196
             7.3
                   1038
                        7
                           ABM88777
                                                    Abm88777 Rice abio
26
    181.5
             6.7
                   564 8
                           ADY23792
                                                    Ady23792 Plant ful
27
    167.5
             6.2
                    767 6 ABR53431
                                                    Abr53431 Protein s
    167.5
28
             6.2
                    767 7 ADK64670
                                                    Adk64670 Disease t
29
    164.5
                    554 3 AAG36165
             6.1
                                                    Aag36165 Arabidops
30
    164.5
                    652 3 AAG36164
             6.1
                                                    Aaq36164 Arabidops
                    781 3 AAG36163
31
    164.5
             6.1
                                                    Aaq36163 Arabidops
                    815 5 AAG78388
32
    162.5
             6.0
                                                    Aag78388 Human H37
33
    162.5
             6.0
                   815 7
                           AAE38620
                                                    Aae38620 Human H37
    162.5
             6.0
                   815 7 AAE38621
34
                                                   Aae38621 Human H37
35
    162.5
             6.0
                    815 8 ADP23184
                                                   Adp23184 PRO polyp
36
    162.5
                                                   Adx05546 Cyclin-de
             6.0
                    815 9 ADX05546
37
      162
             6.0
                    381 2 AAY07056
                                                    Aay07056 Renal can
38
    157.5
             5.8
                        7
                    852
                           ADD45318
                                                    Add45318 Rat Prote
39
    157.5
             5.8
                    852
                        7
                           ADE56352
                                                    Ade56352 Rat Prote
40
    156.5
             5.8
                    573
                           ADM19760
                                                   Adm19760 Protein e
41
      156
             5.8
                    852 7
                           AEA79131
                                                   Aea79131 Human apo
42
      156
             5.8
                    852 9 ADX07612
                                                   Adx07612 Cyclin-de
43
      156
             5.8
                    929 4 AAM78604
                                                   Aam78604 Human pro
44
      156
                    930 8 ABM82400
             5.8
                                                   Abm82400 Tumour-as
                    930 9 ADX07610
45
      156
             5.8
                                                    Adx07610 Cyclin-de
```

# ALIGNMENTS

```
AAU06524
      AAU06524 standard; protein; 504 AA.
XX
AC
     AAU06524;
XX
 DΤ
     24-OCT-2001 (first entry)
XX
      Prostate and testis-related gene 84P2A9 encoded protein.
 DE
XX
 KW
      84P2A9-related protein; prostate; testis; tissue; cancer; leukaemia;
      tumour; kidney; brain; bone; skin; ovary; breast; pancreas; colon; lung;
 KW
. KW
      cytostatic; gene therapy; antibody therapy; ribozyme; serum; blood;
KW
      single chain monoclonal antibody; urine.
XX
```

RESULT 1

```
OS
    Homo sapiens.
XX
PN
    W0200155391-A2.
XX
    02-AUG-2001.
PD
XX
ΡF
    26-JAN-2001; 2001WO-US002651.
XX
    26-JAN-2000; 2000US-0178560P.
PR
XX
PΑ
    (UROG-) UROGENESYS INC.
XX
    Jakobovits A, Afar DEH, Challita-Eid PM, Levin E, Mitchell SC;
PΙ
PΙ
    Hubert RS;
XX
DR
    WPI; 2001-502631/55.
    N-PSDB; AAS11663.
DR
XX
PT
    New 84P2A9 gene and its encoded protein, useful for diagnosing and
    treating cancer, e.g. leukemia and cancer of the prostate, testis,
PT
    kidney, brain or bone, or for eliciting an immune response.
PT
XX
PS
    Claim 13; Fig 2; 149pp; English.
XX
    The polypeptide sequences represent the 84P2A9-related protein and
CC
CC
    peptide fragments of the protein. 84P2A9 exhibits prostate and testis
CC
    specific expression in normal adult tissue, but it is also aberrantly
    expressed in many cancers including leukaemia and tumours of the
CC
    prostate, testis, kidney, brain, bone, skin, ovary, breast, pancreas,
CC
CC
    colon and lung. The 84P2A9 polynucleotide, its related protein and
CC
    peptide fragments and specific PCR primers are therefore useful for
CC
    diagnosing and treating cancer. A vector comprising a polynucleotide
CC
    which encodes a single chain monoclonal antibody, that immunospecifically
    binds to an 84P2A9-related protein, and a ribozyme capable of cleaving a
CC
CC
    polynucleotide having the 84P2A9 coding sequence, are both useful in the
    preparation of a composition for treating a patient with a cancer that
CC
CC
    expresses 84P2A9. The sequences can be used in diagnostic methods to
CC
    monitor the level of 84P2A9 gene products in serum, blood, urine and
CC
    tissue and to thereby detect the presence of cancerous cells
XX
SO
    Sequence 504 AA;
 Query Match
                        100.0%; Score 2694; DB 4; Length 504;
                        100.0%; Pred. No. 3.9e-228;
 Best Local Similarity
 Matches 504; Conservative
                            0; Mismatches
                                               0; Indels
                                                            0; Gaps
                                                                       0;
           1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qγ
             1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Db
Qу
          61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
             61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Db
Qу
         121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
             121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Db
```

```
181 TKNKVKKRKLKIIROGPKIODEGVVLESEETNOTNKDKMECEEOKVSDELMSESDSSSLS 240
Qy
           181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Db
        241 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qу
           Db
        241 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
        301 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
QУ
           301 ILTGSFPLMSHPSRRGFOARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
Db
        361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
           361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Db
        421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Qy
           421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Db
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
           Db
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
RESULT 2
AAB92632
ID
    AAB92632 standard; protein; 528 AA.
XX
AC
    AAB92632;
XX
DT
    26-JUN-2001 (first entry)
XX
    Human protein sequence SEQ ID NO:10938.
DE
XX
KW
    Human; primer; detection; diagnosis; antisense therapy; gene therapy.
XX
os
    Homo sapiens.
XX
PN
    EP1074617-A2.
XX
    07-FEB-2001.
PD
XX
    28-JUL-2000; 2000EP-00116126.
PF
XX
                99JP-00248036.
    29-JUL-1999;
PR
PR
    27-AUG-1999;
                99JP-00300253.
    11-JAN-2000; 2000JP-00118776.
PR
    02-MAY-2000; 2000JP-00183767.
PR
    09-JUN-2000; 2000JP-00241899.
PR
XX
PΑ
    (HELI-) HELIX RES INST.
XX
    Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PΙ
    Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
PΙ
XX
```

DR WPI; 2001-318749/34.

XX

PT Primer sets for synthesizing polynucleotides, particularly the 5602 fullPT length cDNAs defined in the specification, and for the detection and/or
PT diagnosis of the abnormality of the proteins encoded by the full-length
PT cDNAs.

XX

PS Claim 8; SEQ ID NO 10938; 2537pp + Sequence Listing; English.

XX

The present invention describes primer sets for synthesising 5602 fulllength cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention

Sequence 528 AA;

Query Match

CC

XX SO

```
Best Local Similarity
                   100.0%;
                         Pred. No. 4.2e-228;
 Matches 504; Conservative
                      0; Mismatches
                                     0: Indels
                                               0;
                                                  Gaps
                                                        0:
Qу
        1 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 60
          Db
        25 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 84
        61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDOMLVAKRRPSSNLNNNVRGKRPLWH 120
Qу
          85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 144
Db
       121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qу
          145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204
Db
       181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Qу
          205 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 264
Db
       241 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qy
          Db
       265 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 324
```

100.0%; Score 2694; DB 4;

Length 528;

```
301 ILTGSFPLMSHPSRRGFOARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
QУ
            Db
         325 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 384
         361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
            Db
         385 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 444
         421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Qу
            445 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 504
Db
         481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
            Db
         505 GLGLGFPLPKSTSATTTPNAGKSA 528
RESULT 3
ABB97288
    ABB97288 standard; protein; 528 AA.
XX
AC
    ABB97288;
XX
DT
    28-JUN-2002 (first entry)
XX
DΕ
    Novel human protein SEQ ID NO: 556.
XX
KW
    Human; antianaemic; vulnerary; antiinflammatory; immunomodulator;
    antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy;
KW
KW
    neuroprotective; antiparkinsonian; protein therapy; EST;
KW
    expressed sequence tag.
XX
OS
    Homo sapiens.
XX
PN
    W0200222660-A2.
XX
PD
    21-MAR-2002.
XX
ΡF
    10-SEP-2001; 2001WO-US026015.
XX
PR
    11-SEP-2000; 2000US-00659671.
XX
PA
    (HYSE-) HYSEQ INC.
XX
PΙ
    Tang YT, Liu C,
                    Zhou P, Asundi V, Zhang J, Zhao QA, Ren F;
PΙ
    Xue AJ, Yang Y,
                    Wehrman T, Drmanac RT;
XX
DR
    WPI; 2002-292408/33.
    N-PSDB; ABN32474.
DR
XX
    An isolated polynucleotide for treating diseases associated with its
PT
PT
    encoded polypeptide such as cancer and multiple sclerosis.
XX
PS
    Example 2; SEQ ID NO 556; 509pp; English.
XX
    The present invention provides the protein and coding sequences of 444
CC
```

```
CC
    novel human proteins. These were isolated from expressed sequences tags
    (ESTs). They can be used to stimulate cell growth, to regulate
CC
CC
    haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth
CC
    e.g. in burn treatment, to regulate the immune system e.g. to treat
CC
    multiple sclerosis, to regulate activin or inhibin e.g. to treat
CC
    infertility, to regulate haemostasis or thrombolysis e.g. to treat stroke
CC
    and cancer, to screen for drugs, to treat inflammatory conditions e.g.
    rheumatoid arthritis, and to treat nervous system disorders e.g.
CC
    Parkinson's disease. The present sequence is a protein of the invention
CC
XX
SO
    Sequence 528 AA;
                     100.0%; Score 2694; DB 5;
 Query Match
                                            Length 528;
 Best Local Similarity
                    100.0%; Pred. No. 4.2e-228;
 Matches 504; Conservative
                         0; Mismatches
                                         0;
                                            Indels
                                                    0;
                                                        Gaps
                                                              0;
         1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qу
           25 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 84
Db
         61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Qу
           85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDOMLVAKRRPSSNLNNNVRGKRPLWH 144
Db
        121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qv
           145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204
Db
        181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Qу
           205 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 264
Db
        241 STDAGLFTNDEGROGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qу
           265 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 324
Db
        301 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
Qу
           Db
        325 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 384
        361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
           Db
        385 SHHHDHWFSPGARTEHDOHOLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 444
        421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Qу
           Db
        445 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 504
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
           Db
        505 GLGLGFPLPKSTSATTTPNAGKSA 528
```

RESULT 4 ADR99239

ID ADR99239 standard; protein; 376 AA.

XX

```
AC
    ADR99239;
XX
DT
     02-DEC-2004 (first entry)
XX
     Hypothetical protein FLJ10252, SEQ ID 245.
DE
XX
KW
     Cytostatic; breast cancer; cancer; human; FLJ10252.
XX
OS
    Homo sapiens.
XX
PN
    WO2004078035-A2.
XX
     16-SEP-2004.
PD
XX
PF
     27-FEB-2004; 2004WO-US007268.
XX
     28-FEB-2003; 2003US-0450655P.
PR
XX
     (FARB ) BAYER PHARM CORP.
PA
XX
     Eveleigh D, Bigwood D;
PΙ
XX
DR
     WPI; 2004-653556/63.
DR
     N-PSDB; ADR99112.
XX
     Diagnosing breast cancer comprises comparing the level of expression of
PΤ
РΤ
     genes or gene products in a first biological sample taken from a patient
PT
     with that in a normal patient sample.
XX
     Claim 3; SEQ ID NO 245; 53pp; English.
PS
XX
     The present invention relates to a method (M1) for diagnosing breast
CC
CC
     cancer in a patient. The method comprises comparing the level of
     expression of one or more genes or gene products in a biological sample
CC
CC
     from the patient with that in a normal patient sample, where a difference
CC
     in the gene expression in the first sample compared to that in the second
     sample is a diagnostic of the disease. Also claimed are: method (M2) for
CC
     distinguishing between normal and disease tissues; method (M3) for
CC
CC
     monitoring the response of a breast cancer patient to treatment with an
CC
     anti-cancer agent; method (M4) for identifying a compound for treating
CC
     breast cancer; and an array for distinguishing between normal and disease
CC
     tissues comprising two or more probes corresponding to genes selected
CC
     from ADR98995-ADR99121 or comprising two or more polypeptides selected
CC
     from ADR99122-ADR99248. In M1 and M2 the genes are selected from ADR98995
     -ADR99121 and the gene products are polypeptides selected from ADR99122-
CC
     ADR99248. M1 is useful for diagnosing breast cancer. M2 and the array are
CC
CC
     useful for distinguishing between normal and disease tissue. M3 is useful
     for monitoring the response of a breast cancer patient to treatment with
CC
CC
     an anti-cancer agent. M4 is useful for identifying a compound for
     treating breast cancer. Note: The sequence data for this patent did not
CC
     form part of the printed specification, but was obtained in electronic
CC
     format directly from WIPO at ftp.wipo.int/pub/published pct_sequences.
CC
XX
SO
     Sequence 376 AA;
```

67.1%; Score 1808; DB 8; Length 376;

Best Local Similarity 99.7%; Pred. No. 3.1e-150;

Query Match

```
1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qу
           25 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 84
Db
         61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Qу
           85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDOMLVAKRRPSSNLNNNVRGKRPLWH 144
Db
        121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qy
           145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204
Dh
        181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Qy
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KW
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DR
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1; Mismatches

0; Indels

0;

Gaps

0;

Matches 341; Conservative

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    New isolated polynucleotide and encoded polypeptides, useful in
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PT
    responsible for genetic disorders or other traits and to assess
PΤ
PT
    biodiversity.
XX
PS
    Claim 20; SEQ ID NO 38361; 103pp; English.
XX
CC
    The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC
    sequences. (I) is useful as hybridisation probes, polymerase chain
CC
    reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
CC
    and in recombinant production of (II). The polynucleotides are also used
CC
    in diagnostics as expressed sequence tags for identifying expressed
CC
    genes. (I) is useful in gene therapy techniques to restore normal
    activity of (II) or to treat disease states involving (II). (II) is
CC
    useful for generating antibodies against it, detecting or quantitating a
CC
    polypeptide in tissue, as molecular weight markers and as a food
CC
    supplement. (II) and its binding partners are useful in medical imaging
CC
    of sites expressing (II). (I) and (II) are useful for treating disorders
CC
    involving aberrant protein expression or biological activity. The
CC
    polypeptide and polynucleotide sequences have applications in
CC
    diagnostics, forensics, gene mapping, identification of mutations
CC
    responsible for genetic disorders or other traits to assess biodiversity
CC
    and to produce other types of data and products dependent on DNA and
CC
    amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic
CC
    amino acid sequences of the invention. Note: The sequence data for this
CC
    patent did not appear in the printed specification, but was obtained in
CC
    electronic format directly from WIPO at
CC
CC
    ftp.wipo.int/pub/published pct sequences
XX
    Sequence 313 AA;
SQ
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## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame plus p2n model

Run on: December 11, 2005, 17:36:59; Search time 7575 Seconds

(without alignments)

3782.059 Million cell updates/sec

US-09-771-312-2 Title:

Perfect score: 2694

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5883141 seqs, 28421725653 residues Searched:

Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 2: gb in:\*
- 3: gb env:\*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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AK001114 Homo sapi
AX405697 Sequence
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  AUTHORS
            Ota, T., Isogai, T., Nishikawa, T., Hayashi, K., Saito, K.,
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            PΙ
                 SAITO,
            PΙ
                 JUNICHI YAMAMOTO, SHIZUKO ISHII, TOMOYASU SUGIYAMA, AI
WAKAMATSU,
            PΙ
                 KEIICHI NAGAI, TETSUJI OTSUKI
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Mismatches:

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Percent Similarity:

Best Local Similarity: 100.00%

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5.58e-138

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100.00%

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US-09-771-312-2 (1-504) x BD155908 (1-2338)

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2003
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VERSION
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REFERENCE
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           Ota, T., Isogai, T., Nishikawa, T., Hayashi, K., Saito, K.,
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REFERENCE
 AUTHORS
           Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R.,
           Wakamatsu, A., Hayashi, K., Sato, H., Nagai, K., Kimura, K., Makita, H.,
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  TITLE
            Complete sequencing and characterization of 21,243 full-length
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            Nat. Genet. 36 (1), 40-45 (2004)
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            Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y.,
 AUTHORS
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  TITLE
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            Isogai, T. and Otsuki, T.
  TITLE
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  JOURNAL
            Submitted (16-FEB-2000) Takao Isogai, Helix Research Institute,
            Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
            (E-mail:genomics@hri.co.jp, Tel:81-438-52-3975, Fax:81-438-52-
3986)
COMMENT
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# ORIGIN

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REFERENCE
          Tang, Y.T., Liu, C., Zhou, P., Asundi, V., Zhang, J., Zhao, Q.A.,
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Ren, F.,
          Xue, A.J., Yang, Y., Wehrman, T. and Drmanac, R.T.
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 JOURNAL
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REFERENCE
                (bases 1 to 4537)
  AUTHORS
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M.,
Schuler, G.D.,
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            Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
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            Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
            Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,
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  TITLE
            Generation and initial analysis of more than 15,000 full-length
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            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
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            Strausberg, R.
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  JOURNAL
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            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            NIH-MGC Project URL: http://mgc.nci.nih.gov
  REMARK
COMMENT
            Contact: MGC help desk
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Dr. James Lin, University of Iowa
            cDNA Library Preparation: M. Bento Soares, University of Iowa
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: University of Iowa, Dr. M. Bento Soares and Dr.
            Thomas L. Casavant.
            Web site: http://genome.uiowa.edu
            Contact: bento-soares@uiowa.edu; tom-casavant@uiowa.edu
            Bonaldo, M. F., Akabogu, I., Bair, T., Bair, J., Crouch, K., Davis, A.,
            Fishler, K., Keppel, C., Kucaba, T., Lebeck, M., Melo, A., Schaefer, K.,
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            Clone distribution: MGC clone distribution information can be
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Location/Qualifiers

SOURCE

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US-09-771-312-2 (1-504) x BC054810 (1-4537)
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111

Db

1655 GGAAACCCTGCC 1666

# us-09-771-312-2.rng

### GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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geneseqn2003bs:\*

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9: 10:

11: 12:

13: 14:

## SUMMARIES

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ΡI
PΙ
XX
         WPI: 2001-318749/34.
DR
XX
         Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length
PT
PT
PT
PT
          CDNĂs.
XX
          Claim 8; SEQ ID NO 10937; 2537pp + Sequence Listing; English.
PS
XX
         The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602
CC
CC
CC
CC
         nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC
CC
         of an oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the
CC
\mathsf{CC}
CC
CC
CC
CC
         specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH18742 represent human cDNA segmences: AAB02446 to AAB05803
CC
CC
CC
CC
CC
CC
          AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893
CC
CC
          represent human amino acid sequences; and AAH13629 to AAH13632 represent
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CC
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CC
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us-09-771-312-2.rng
DB: 4 Gaps: 0

US-09-771-312-2 (1-504) x AAH13916 (1-2338) 1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20 Qy 171 ATGGAGGAGCTGGTTCATGACCTTGTCTCAGCATTGGAAGAGCGCAGGCAAGCTCGA 230 Db 21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40 Qy 231 GGTGGATTTGCTGAAACAGGAGACCATTCTCGAAGTATATCTTGCCCTCTGAAACGCCAG 290 Db 41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60 Qy 291 GCÁÁGGÁÁÁÁÁGGÁGÁGÁGÁGÁÁÁÁÁCGGÁGGTCGTÁTÁÁTGTGCÁTCÁCCCGTGGGÁGÁCT 350 Db 61 GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg 80 Qy Db 81 GluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMetLeuValAla 100 Qy 411 GAGAATCACAATAATAATAAAAAAGATCACAGTGACTCTGATGACCAAATGTTAGTAGCA 470 Db 101 LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis 120 Qy 471 ÁÁGCGCÁGGCCGTCÁTCÁÁÁCTTÁÁÁTÁÁTÁÁTGTTCGAGGGAAAAGÁCCTCTÁTGGCÁT 530 Db 121 GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal 140 Qy 531 GAGTCTGATTTTGCTGTGGACAATGTTGGGAATAGAACTCTGCGCAGGAGGAGAAAGGTA 590 Db Qy Db 161 ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe 180 Qy 651 CCTGÁGGGTTGTAGAGATCAGGÁCÁTGGÁCÁGTGATAGAGCCTÁCCAGTATCAAGAÁTTT 710 Db Qy Db 201 AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu 220 Qy 771 GATGAAGGAGTAGTTTTAGAAAGTGAGGAAACGAACCAGACCAGTAAGGACAAAATGGAA 830 Db CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer 240 Qу 831 TGTGAAGAGCAAAAAGTCTCAGATGAGCTCATGAGTGAAAGTGATTCCAGCAGTCTCAGC 890 Db 241 SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer 260 Qy 891 ÁGCÁCTGATGCTGGÁTTGTTTÁCCAATGATGAGGGAAGACAAGGTGATGATGAACAGAGT 950 Db 261 AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp 280 Qy 951 GACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTTGTGCCCTGG 1010 Dh 281 TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer 300 Qy Db 1011 TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT 1070

```
301 IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg 320
Qy
Db
        1071 ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA 1130
         321 LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr 340
Qy
Db
        1131 CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT 1190
         341 SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp 360
Qy
        1191 TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT 1250
Dh
         361 SerHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln 380
Qy
Db
        1251 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1310
         381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
        1311 CTTCTGAGAGATAATCGAGCTGAAAGAGACACAAGAAAAATTGTTCTGTGAGAACAGCC 1370
Db
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
Db
        1371 ÁGCÁGGCÁÁACÁÁGCÁTGCÁTTTÁGGÁTCCTTÁTGCÁCGGGÁGÁTÁTCÁÁACGGÁGÁGÁGÁ 1430
         421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
             1431 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1490
Db
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
             Db
        1491 ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA 1550
         461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
Db
        1551 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1610
         481 GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
             Db
        1611 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1670
         501 GlyLysSerAla 504
Qy
        1671 GGAAAATCCGCC 1682
Db
RESULT 2
ADR99112
    ADR99112 standard; DNA; 2338 BP.
ID
XX
AC
    ADR99112;
XX
    02-DEC-2004 (first entry)
DT
XX
    Hypothetical protein FLJ10252, coding sequence, SEQ ID 118.
DE
XX
    Cytostatic; breast cancer; cancer; human; gene; ds; FLJ10252.
KW
XX
os
    Homo sapiens.
XX
    WO2004078035-A2.
PN
XX
    16-SEP-2004.
PD
XX
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us-09-771-312-2.rna
PF
       27-FEB-2004; 2004WO-US007268.
XX
PR
       28-FEB-2003: 2003US-0450655P.
XX
       (FARB ) BAYER PHARM CORP.
PA
XX
PΙ
       Eveleigh D, Bigwood D;
XX
       WPI; 2004-653556/63.
DR
       P-PSDB; ADR99239.
REFSEQ; NM_018040.1.
DR
DR
XX
       Diagnosing breast cancer comprises comparing the level of expression of genes or gene products in a first biological sample taken from a patient
PT
PT
PT
       with that in a normal patient sample.
XX
PS
       Claim 2; SEQ ID NO 118; 53pp; English.
XX
       The present invention relates to a method (M1) for diagnosing breast
CC
       cancer in a patient. The method comprises comparing the level of expression of one or more genes or gene products in a biological sample
CC
CC
       from the patient with that in a normal patient sample, where a difference
CC
CC
       in the gene expression in the first sample compared to that in the second
       sample is a diagnostic of the disease. Also claimed are: method (M2) for
CC
       distinguishing between normal and disease tissues; method (M3) for
CC
       monitoring the response of a breast cancer patient to treatment with an anti-cancer agent; method (M4) for identifying a compound for treating breast cancer; and an array for distinguishing between normal and disease tissues comprising two or more probes corresponding to genes selected from ADR98995-ADR99121 or comprising two or more polypeptides selected from ADR98122-ADR99248. In M1 and M2 the genes are selected from ADR98995
CC
CC
CC
CC
CC
CC
       -ADR99121 and the gene products are polypeptides selected from ADR99122-ADR99248. M1 is useful for diagnosing breast cancer. M2 and the array are useful for distinguishing between normal and disease tissue. M3 is useful for monitoring the response of a breast cancer patient to treatment with an anti-cancer agent. M4 is useful for identifying a compound for
CC
CC
CC
CC
CC
       treating breast cancer. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic
CC
CC
       format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.
CC
XX
       Sequence 2338 BP; 739 A; 478 C; 541 G; 580 T; 0 U; 0 Other;
SO
Alignment Scores:
Pred. No.:
                                   2.07e-199
                                                                               2338
                                                         Length:
Score:
                                   2694.00
                                                         Matches:
                                                                               504
Percent Similarity:
                                   100.00%
                                                         Conservative:
                                                                               0
Best Local Similarity:
                                   100.00%
                                                         Mismatches:
                                                                               0
                                                         Indels:
Query Match:
                                   100.00%
                                                                               0
                                                                               0
                                                         Gaps:
US-09-771-312-2 (1-504) x ADR99112 (1-2338)
Qy
                 1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20
                     Db
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                21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
Db
              231 GGTGGATTTGCTGAAACAGGAGACCATTCTCGAAGTATATCTTGCCCTCTGAAACGCCAG 290
                41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
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291 GCAAGGAAAAGGAGAGGAGAAAACGGAGGTCGTATAATGTGCATCACCCGTGGGAGACT 350

Db

Qy	61	GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg	80
Db	351	GGTCACTGCTTAAGTGAAGGCTCTGATTCTAGTTTAGAAGAACCAAGCAAG	410
Qy	81	GluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMetLeuValAla	100
Db	411	ĠĠĠĠĂĊŢĊĠĊĠĠŢĠĠĊĊĠĠŢĠĠĊĊĊĠĠŢĠĠĊĊĠĠĠŢĠĠĊĠĠĠĠĠĠ	470
Qy	101	LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis	120
Db		AAGCGCAGGCCGTCATCAAACTTAAATAATGTTCGAGGGAAAAGACCTCTATGGCAT	530
Qy	121	GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal	140
Db		GAGTCTGATTTTGCTGTGGACAATGTTGGGAATAGAACTCTGCGCAGGAGGAGAAAGGTA	
Qy		LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro	
Db		AAACGCATGGCAGTAGATCTCCCACAGGACATCTCTAACAAACGGACAATGACCCAGCCA	
Qy Dh		ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe	
Db		CCTGAGGGTTGTAGAGATCAGGACATGGACAGTGATAGAGCCTACCAGTATCAAGAATTT ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln	710
Qy Db			
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Db			
Qy		CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	
Db			
Qy	241	SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	260
Db	891	AGCACTGATGCTGGATTGTTTACCAATGATGAGGGAAGACAAGGTGATGATGAACAGAGT	950
Qy	261	AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	280
Db	951		1010
Qy	281	TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	300
Db		TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT	1070
Qy	301	<pre>IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg</pre>	320
Db	1071	ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA	1130
Qy	321	LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	340
Db	1131	CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAAATCTGGAGGGACTCCAACT	1190
Qy	341	SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	360
Db		TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTCCCCGGAT	1250
Qy	361	SerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	380

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us-09-771-312-2.rng
Db
        1251 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1310
         381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
        1311 CTTCTGÁGÁGÁTÁÁTCGÁGCTGAAAGÁGÁCÁCAÁGÁÁÁAATTGTTCTGTGÁGÁÁÁÁCAGCC 1370
Db
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
             Db
        1371 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA 1430
         421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
             Db
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         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
             Db
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         461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
Db
        1551 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1610
         481 GlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
             Db
        1611 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1670
         501 GlyLysSerAla 504
Qy
             Db
        1671 GGAAAATCCGCC 1682
RESULT 3
ABN59701
    ABN59701 standard; cDNA; 2344 BP.
ΙD
XX
    ABN59701:
AC
XX
    28-JUN-2002 (first entry)
DT
XX
    Novel human coding sequence SEQ ID NO: 112.
DE
XX
    Human; antianaemic; vulnerary; antiinflammatory; immunomodulator; antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy; neuroprotective; antiparkinsonian; protein therapy; EST;
KW
KW
KW
    expressed sequence tag; gene; ss.
KW
XX
os
    Homo sapiens.
XX
PΝ
    WO200222660-A2.
XX
PD
    21-MAR-2002.
XX
    10-SEP-2001; 2001WO-US026015.
PF
XX
    11-SEP-2000; 2000US-00659671.
PR
XX
PA
    (HYSE-) HYSEQ INC.
XX
ΡI
    Tang YT, Liu C, Zhou P, Asundi V, Zhang J, Zhao QA,
                                                           Ren F;
    Xue AJ, Yang Y, Wehrman T, Drmanac RT;
PΙ
XX
    WPI; 2002-292408/33.
DR
    P-PSDB; ABB97288.
DR
XX
```

```
us-09-771-312-2.rng
PT
      An isolated polynucleotide for treating diseases associated with its
      encoded polypeptide such as cancer and multiple sclerosis.
PT
XX
      Claim 1; SEQ ID NO 112; 509pp; English.
PS
XX
CC
      The present invention provides the protein and coding sequences of 444
     The present invention provides the protein and coding sequences of 444 novel human proteins. These were isolated from expressed sequences tags (ESTs). They can be used to stimulate cell growth, to regulate haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth e.g. in burn treatment, to regulate the immune system e.g. to treat multiple sclerosis, to regulate activin or inhibin e.g. to treat infertility, to regulate haemostasis or thrombolysis e.g. to treat stroke and cancer, to screen for drugs, to treat inflammatory conditions e.g. rheumatoid arthritis, and to treat nervous system disorders e.g. Parkinson's disease. The present sequence is a coding sequence of the invention
CC
CC
CC
CC
CC
CC
CC
CC
CC
      invention
CC
XX
     Sequence 2344 BP; 747 A; 476 C; 541 G; 580 T; 0 U; 0 Other;
SQ
Alignment Scores:
Pred. No.:
                            2.08e-199
                                                               2344
                                              Length:
                            2694.00
                                                               504
Score:
                                              Matches:
Percent Similarity:
                            100.00%
                                              Conservative:
                                                               0
Best Local Similarity:
                            100.00%
                                              Mismatches:
                                                               0
Query Match:
                            100.00%
                                              Indels:
                                                               0
                                              Gaps:
                                                               0
US-09-771-312-2 (1-504) x ABN59701 (1-2344)
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            21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
           230 GGTGGÁTTTGCTGÁÁÁCÁGGÁGÁCCÁTTCTCGÁÁGTÁTATCTTGCCCTCTGÁÁÁCGCCÁG 289
Db
             41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
           290 GCAAGGAAAAGGAGAGGAGAAAACGGAGGTCGTATAATGTGCATCACCCGTGGGAGACT 349
Db
            61 GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg 80
Qy
Db
           Qy
Db
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Qy
           470 AAGCGCAGGCCGTCATCAAACTTAAATAATAATGTTCGAGGGAAAAGACCTCTATGGCAT 529
Db
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Qу
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Db
Qy
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                Db
           590 AAACGCATGGCAGTAGATCTCCCACAGGACATCTCTAACAAACGGACAATGACCCAGCCA 649
           161 ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe 180
Qy
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Db		ACCAAGAACAAAGTCAAAAAAAGAAAGTTGAAAATAATCAGACAAGGACCAAAAATCCAA	
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Db		GATGAAGGAGTAGTTTTAGAAAGTGAGGAAACGAACCAGACCAATAAGGACAAAATGGAA CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	
Qy Db			889
Qy		SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	
Db			
Qy	261	AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	280
Db	950		1009
Qy	281	TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	300
Db	1010	TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT	1069
Qy			320
Db		ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA	1129
Qy Db		LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	340 1189
Qy		SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	
Db			1249
Qy	361	SerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	380
Db	1250	TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG	1309
Qy	381	LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla	400
Db		CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAAATTGTTCTGTGAGAACAGCC	
Qy		SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg	
Db		AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA	
Qy Db		LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro	
Qy		IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer	
Db			
Qy	461	GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys	480
Db	1550		1609
Qy	481	GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla	500

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us-09-771-312-2.rng
                          Db
                1610 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1669
                  501 GlyLysSerAla 504
Qy
                          Db
                1670 GGAAAATCCGCC 1681
RESULT 4
AAS11663
         AAS11663 standard; cDNA; 2345 BP.
ID
XX
AC
         AAS11663;
XX
         24-0CT-2001 (first entry)
DT
XX
DE
         Prostate and testis-related gene 84P2A9 cDNA.
XX
         84P2A9; PCR primer; DNA adaptor; prostate; testis; tissue; cancer; ss; leukaemia; tumour; kidney; brain; bone; skin; ovary; breast; pancreas;
KW
KW
         colon; lung; cytostatic; gene therapy; antibody therapy; ribozyme; single chain monoclonal antibody; serum; blood; urine.
KW
KW
XX
         Homo sapiens.
05
XX
PN
         WO200155391-A2.
XX
         02-AUG-2001.
PD
XX
PF
         26-JAN-2001; 2001WO-US002651.
XX
PR
         26-JAN-2000; 2000US-0178560P.
XX
PA
         (UROG-) UROGENESYS INC.
XX
PΙ
         Jakobovits A, Afar DEH, Challita-Eid PM, Levin E, Mitchell SC;
ΡI
         Hubert RS:
XX
         WPI: 2001-502631/55.
DR
DR
         P-PSDB; AAU06524.
XX
PT
         New 84P2A9 gene and its encoded protein, useful for diagnosing and
         treating cancer, e.g. leukemia and cancer of the prostate, testis, kidney, brain or bone, or for eliciting an immune response.
PT
PT
XX
PS
         Claim 1; Fig 2; 149pp; English.
XX
        The nucleic acid sequences represent the 84P2A9 gene and the primers and adaptors used to amplify 84P2A9 DNA. 84P2A9 exhibits prostate and testis specific expression in normal adult tissue, but it is also aberrantly expressed in many cancers including leukaemia and tumours of the prostate, testis, kidney, brain, bone, skin, ovary, breast, pancreas, colon and lung. The 84P2A9 polynucleotide, its related protein and also peptide fragments of the protein are therefore useful for diagnosing and
CC
CC
CC
CC
CC
CC
CC
        treating cancer. A vector comprising a polynucleotide which encodes a single chain monoclonal antibody, that immunospecifically binds to an 84P2A9-related protein, and a ribozyme capable of cleaving a polynucleotide having the 84P2A9 coding sequence, are both useful in the preparation of a composition for treating a patient with a cancer that expresses 84P2A9. The sequences can be used in diagnostic methods to monitor the level of 84P2A9 gene products in serum, blood, urine and tissue and to thereby detect the presence of cancerous cells
CC
CC
CC
CC
CC
CC
CC
CC
         tissue and to thereby detect the presence of cancerous cells
XX
SQ
         Sequence 2345 BP; 750 A; 476 C; 542 G; 577 T; 0 U; 0 Other;
```

Alignment Scores: Pred. No.: Score: Percent Similarity: Best Local Similarity: Query Match: DB:			2.08e-199 2694.00 100.00% 100.00% 100.00%	Length: Matches: Conservative: Mismatches: Indels: Gaps:	2345 504 0 0 0
US-09-771	-312	-2 (1-504)	x AAS11663 (1-	-2345)	
Qy	1	MetGluGlu	LeuValHisAspLeı	uValSerAlaLeuGlu	GluSerSerGluGlnAlaArg 20 
Db	163	ATGGAGGAG	CTGGTTCATGACCT	TGTCTCAGCATTGGAA	GAGAGCTCAGAGCAAGCTCGA 222
Qy	21	GlyGlyPhe	AlaGluThrGlyAsı	oHisSerArgSerIle	SerCysProLeuLysArgGln 40 
Db	223	GGTGGATTT	GCTGAAACAGGAGAG	CCATTCTCGAAGTATA	TCTTGCCCTCTGAAACGCCAG 282
Qy	41	AlaArgLys,	ArgArgGlyArgLy:	sArgArgSerTyrAsn	ValHisHisProTrpGluThr 60 
Db	283				GTGCATCACCCGTGGGAGACT 342
Qy	61	GlyHisCys	LeuSerGluGlySe	rAspSerSerLeuGlu 	GluProSerLysAspTyrArg 80 
Db	343	ĠĠŦĊĂĊŦĠĊ	ŤŤÁÁĠŤĠÁÁĠĠĊŤĊ	ÍGÁTTCTÁGTTTÁGÁÁ	GAACCAAGCAAGGACTATAGA 402
Qy	81	GluAsnHis,	AsnAsnAsnLysLy: 	sAspHisSerAspSer 	AspAspGlnMetLeuValAla 100 
Db					GATGACCAAATGTTAGTAGCA 462
Qy					GlyLysArgProLeuTrpHis 120
Db					GGGAAAAGACCTCTATGGCAT 522
Qy				111111111	LeuArgArgArgArgLysVal 140
Db					CTGCGCAGGAGGAGAAGGTA 582
Qy					LysArgThrMetThrGlnPro 160
Db					AAACGGACAATGACCCAGCCA 642 AlatyrGlnTyrGlnGluPhe 180
Qy Db			1) [[[[]] []] [[] [[]]		
Qy					ArgGlnGlyProLysIleGln 200
Db					
Qy					ThrAsnLysAspLysMetGlu 220
Db					
Qy	221	CysGluGlu	GlnLysValSerAsı	oGluLeuMetSerGlu	SerAspSerSerSerLeuSer 240
Db			111111111111		
Qy	241	SerThrAsp	A]aG]yLeuPheThi	ŗĄṣṇĄṣpĢļuĢļyĄrg	G]nG]yAspAspG]uG]nSer 260
Db	883	AGCACTGAT			

```
us-09-771-312-2.rng
         261 AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp 280
Qy
         943 GACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTTGTGCCCTGG 1002
Db
         281 TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer 300
Qy
        1003 TGGGÁAÁÁGGÁÁGÁTCCTÁCTGÁGCTÁGÁCÁÁÁÁÁTGTÁCCÁGÁTCCTGTCTTTGÁÁÁGT 1062
Db
         301 IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg 320
Qy
            1063 ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA 1122
Db
         321 LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr 340
Qy
Db
        1123 CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT 1182
         341 SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp 360
Qy
Db
        1183 TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT 1242
         361 SerHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln 380
Qy
Db
        1243 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1302
         381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
Db
        1303 CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAGAAAAATTGTTCTGTGAGAACAGCC 1362
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
             1363 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA 1422
Db
         421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
Db
        1423 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1482
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
            Db
        1483 ATCCTÁGÁÁÁÁTÁÁTATTGGÁÁÁÁCCGÁÁTGCTTCÁGÁÁTÁTGGGCTGGÁCGCCTGGGTCÁ 1542
        Qy
Db
         481 GlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qγ
Db
        1603 GGÁTTÁGGÁCTTGGÁTTTCCTCTÁCCAÁAÁÁGTACTTCCGCÁÁCTÁCTACCCCCAATGCÁ 1662
        501 GlyLysSerAla 504
Qy
Db
        1663 GGAAAATCCGCC 1674
RESULT 5
ACN91982
    ACN91982 standard; DNA; 2583 BP.
ID
XX
AC
    ACN91982;
XX
    02-DEC-2004 (first entry)
DT
XX
    Breast cancer related marker, seq id 13132.
DE
XX
    Cancer; breast; tumour; cytostatic; marker; detection; therapy; ds.
KW
```

```
XX
      Homo sapiens.
os
XX
PN
      US2003099974-A1.
XX
PD
      29-MAY-2003.
XX
      18-JUL-2002; 2002US-00198846.
PF
XX
PR
      18-JUL-2001; 2001US-0306220P.
XX
PA
      (MILL-) MILLENNIUM PHARM INC.
XX
PΙ
      Lillie J, Xu Y, Wang Y, Steinmann K;
XX
     WPI: 2003-787014/74.
DR
XX
      Novel isolated polypeptide associated with breast cancer, useful for
PT
PT
      detecting presence of polypeptide in sample, as a marker for breast
PT
      cancer.
XX
PS
      Disclosure; SEQ ID NO 13132; 36pp; English.
XX
CC
      The invention relates to an isolated polypeptide (I) associated with
     breast cancer which is encoded by a nucleic acid molecule comprising a nucleotide sequence (S1). Further disclosed is an antibody that binds to the polypeptide of the invention. The activity of the polypeptide of the invention may be described as cytostatic. The antibody is useful for detecting the presence of (I) in a sample. Nucleic acid molecules of the
CC
CC
CC
CC
CC
      invention are useful in the detection of breast tumours. (I) is useful as
CC
      a marker for breast cancer and in breast cancer therapy. Sequences given
CC
      in records ACN78851-ACN92934 represent nucleic acid markers associated
CC
     with breast cancer. Note: The sequence listing does not form part of the specification but may be obtained in electronic format from the USPTO web site at seqdata.uspto.gov/sequence.html?DocID=20030099974
CC
CC
CC
XX
      Sequence 2583 BP; 813 A; 519 C; 575 G; 664 T; 0 U; 12 Other:
SQ
Alignment Scores:
Pred. No.:
                            2.35e-199
                                             Lenath:
                                                              2583
                            2694.00
Score:
                                             Matches:
                                                              504
Percent Similarity:
                            100.00%
                                             Conservative:
                                                              0
Best Local Similarity:
                            100.00%
                                             Mismatches:
                                                              0
Query Match:
                            100.00%
                                             Indels:
                                                              0
                                                              0
DB:
                            11
                                             Gaps:
US-09-771-312-2 (1-504) x ACN91982 (1-2583)
Qy
              1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20
                Db
           199 ATGGAGGAGCTGGTTCATGACCTTGTCTCAGCATTGGAAGAGCTCAGAGCAAGCTCGA 258
            21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
                 259 GGTGGATTTGCTGAAACAGGAGACCATTCTCGAAGTATATCTTGCCCTCTGAAACGCCAG 318
Db
            41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
Db
           319 GCAAGGAAAAGGAGAGGAGAAAACGGAGGTCGTATAATGTGCATCACCCGTGGGAGACT 378
            61 GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg 80
Qy
           Db
```

Qy	81	GluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMetLeuValAla	100
Db	439		498
Qy	101	LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis	120
Db	499	AAGCGCAGGCCGTCATCAAACTTAAATAATAATGTTCGAGGGAAAAGACCTCTATGGCAT	558
Qy	121	GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal	140
Db	559	GAGTCTGATTTTGCTGTGGACAATGTTGGGAATAGAACTCTGCGCAGGAGGAGAAAGGTA	618
Qy	141	LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro	160
Db	619	AAACGCATGGCAGTAGATCTCCCACAGGACATCTCTAACAAACGGACAATGACCCAGCCA	678
Qy	161	ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe	180
Db	679	ĊĊŦĠĂĠĠĠŦŦĠŦĂĠĂĠĂŦĊĂĠĠĂĊĂŦĠĠĂĊĀĠŦĠĀŦĀĠĀĠĊĊŦĂĊĊĀĠŦĂŦĊĀĀĠĀĀŦŦŤ	738
Qy	181	ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln	200
Db		ACCAAGAACAAAGTCAAAAAAAGAAAGTTGAAAATAATCAGACAAGGACCAAAAATCCAA	
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Db		ĠÀŤĠÀĠĠĀĠŤĀĠŤŤŤŤĀĠÀÁĀĠŤĠĀĠĠĀĀĀĊĠAACĊĀĠĀCCAATAAGGACAAAATGGAA	
Qy	221	CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	240
Db		TGTGAAGAGCAAAAAGTCTCAGATGAGCTCATGAGTGAAAGTGATTCCAGCAGTCTCAGC	
Qy		SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	
Db		AGCACTGATGCTGGATTGTTTACCAATGATGAGGGAAGACAAGGTGATGATGAACAGAGT	
Qy		AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	
Db		GACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTTGTGCCCTGG	
Qy		TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	
Db		TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT	
Qy		<pre>IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg                                     </pre>	
Db		ÁTCTTÁACTGGTTCTTTTCCCCTTÁTGTCÁCÁCCCÁÁGCÁGÁAGÁGAGGTTTCCÁÁGCTÁGÁ LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	
Qy Db			
Db		SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	
Qy Db		TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT	
Db		SerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	
Qy Db		TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG	
		LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla	
Qy	301	LeuLeuArgAspAshArgAraGruArgGryArsLysLysAshCys3ervarArgrin Ara	700

```
us-09-771-312-2.rng
        1339 CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAATTGTTCTGTGAGAACAGCC 1398
Db
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
Db
         1399 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA 1458
          421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
              1459 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1518
Db
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
Db
         1519 ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA 1578
          461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
              1579 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1638
Db
         481 GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
Db
         1639 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1698
          501 GlyLysSerAla 504
Qy
              1699 GGAAAATCCGCC 1710
Db
RESULT 6
AAS72189
     AAS72189 standard; cDNA; 1563 BP.
ID
XX
     AAS72189;
AC
XX
     13-FEB-2002 (first entry)
DT
XX
     DNA encoding novel human diagnostic protein #7993.
DE
XX
     Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW
     food supplement; medical imaging; diagnostic; genetic disorder; ss.
KW
XX
     Homo sapiens.
os
XX
    WO200175067-A2.
PΝ
XX
PD
     11-oct-2001.
XX
     30-MAR-2001; 2001WO-US008631.
PF
XX
     31-MAR-2000; 2000US-00540217.
23-AUG-2000; 2000US-00649167.
PR
PR
XX
     (HYSE-) HYSEQ INC.
PA
XX
PΙ
     Drmanac RT, Liu C, Tang YT;
XX
     WPI: 2001-639362/73.
DR
     P-PSDB; ABG08002.
DR
XX
     New isolated polynucleotide and encoded polypeptides, useful in
PT
     diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess
PT
PT
     biodiversity.
PT
XX
     Claim 1; SEQ ID NO 7993; 103pp; English.
PS
```

```
XX
       The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal activity of (II) or to treat disease states involving (II). (II) is
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CC
CC
CC
        useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and
CC
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CC
CC
        and to produce other types of data and products dependent on DNA and amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention. Note: The sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPO at
CC
CC
cc
CC
CC
        ftp.wipo.int/pub/published_pct_sequences
CC
XX
        Sequence 1563 BP; 486 A; 326 C; 395 G; 356 T; 0 U; 0 Other;
SQ
Alignment Scores:
Pred. No.:
                                        1.55e-106
1495.50
                                                                 Length:
                                                                                          1563
Score:
                                                                 Matches:
                                                                                          312
Percent Similarity:
                                        69.80%
                                                                 Conservative:
                                        68.27%
Best Local Similarity:
                                                                 Mismatches:
                                                                                          21
Query Match:
                                        55.51%
                                                                 Indels:
                                                                                          117
                                                                 Gaps:
US-09-771-312-2 (1-504) x AAS72189 (1-1563)
                    1 MetGluGluLeuValHisAspLeuValSerAlaLeuGlu---GluSerSerGluGlnAla 19
Qy
                155 ÁTGGÁGGÁGCTGGTTCÁTGÁCCTTGTCTCÁGCÁTTGGÁÁAGAGÁGCTCCAGAGCAAGCCT 214
Dh
                Qy
Db
                  39 rgGlnAlaArgLysArgArgGlyArgLysArgArg-SerTyrAsnValHisHisProTrp 58
Qy
                275 GCCCAGCAAGGAAAAGGAGAGGGAGAAAACGGAGGTTCGTATAATGTGCATCACCCGTGG 334
Db
                  59 Glu-ThrGlyHisCysLeu--SerGluGlySerAspSerSerLeuGluGluProSerLys 77
Qy
                335 GÁGGÁCTGGTCÁCTGGCTTAAAGTGÁAGGCTCTGATTCTAGT------ 376
Db
Qy
                  78 AspTyrArgGluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMet 97
                376 ----- 376
Db
                  98 LeuValAlaLysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgPro 117
Qy
                376 ----- 376
Db
                118 LeuTrpHisGluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArg 137
Qy
                376 ----- 376
Db
```

		us-09-771-312-2.rng	
Qy		ArgLysValLysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMet	
Db	376		376
Qy		ThrGlnProProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyr	
Db	376		376
Qy		GlnGluPheThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyPro	
Db	377	TTTAGAAGAACAÀAĠTCAÀAÀAÀAÀAĠAÁĠTTĠAAAATAATĊAĠAĊAAĠĠAĊĊA	430
Qy	198	LysIleGlnAspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAsp	217
Db	431	ÁÁÁÁTCCÁÁGÁTGÁÁGGÁGTÁGTTTTÁGÁÁÁÁGTGÁGGÁÁACGÁÁCCÁGÁCCÁÁTÁÁGGÁC	490
Qy	218	LysMetGluCysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSer	237
Db	491	ÁÁÁÁTGGÁÁTGTGÁÁGÁGCÁÁÁÁÁÁGTCTCÁGÁTGÁGCTCÁTGÁGTGÁÁÁÁGTGÁTTCCÁGC	550
Qy		SerLeuSerSerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAsp	
Db		AGTCTCAGCAGCACTGATGCTGGATTGTTTACCAATGATGAGGGAAGACAAGGTGATGAT	
Qy		GluGlnSerAspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyVal	
Db		GAACAGAGTGACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTT	
Qy		ValProTrpTrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProVal	
Db		GTGCCCTGGTGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTC	
Qy		PheGluSerIleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGly-Ph	317
Db		TTTGAAAGTATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTT	790
Qy	317	eGlnAlaArgLeuSerArg-LeuHisGlyMetSerSerLysAsnIleLysLysSerGlyG	337
Db		CCAACTAAGACTCAGTCGGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAG	
Qy o'		lyThrProThrSerMetValProIleProGlyProValGlyAsnLysArgMetValHisP	
Db		GGACTCCAACTTCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATT	
Qy Di-		heSerProAspSerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisA	
Db		TTTCCCCGGATTCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATG	
Qy Dh		spGlnHisGlnLeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerV	
Db		ACCAGCATCAGCTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAAATTGTTCTG	
Qy Dh		alArgThrAlaSerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleL	
Db		TGAGAACAGCCAGCAGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCA	
Qy Db		ysArgArgArgLysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluA	
		AACGGAGAAAAGCTGCACCTTTGCCTGGACCTACTACTGCAGATTATTTCTCCCCCA	TT20
Qy Db		snAlaGlnProIleLeuGluAsnAsnIleGlyAsn 448 :::   :::: TTCCCAAGCCAGTTATAGTAAAAGAATGTGGAAGT 1185	
UU	TTJT	IICCCAAGCCAGIIAIAGIAAAAGAAIGIGGAAGI 1180	

### GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: December 4, 2005, 10:07:40; Search time 41 Seconds

(without alignments)

1182.762 Million cell updates/sec

Title: US-09-771-312-2

Perfect score: 2694

Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5.

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

0.

Maximum Match 100%

Listing first 45 summaries

Database: PIR 80:\*

1: pir1:\*

2: pir2:\*

3: pir3:\*

4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result		∛ Query				
No.	Score		Length	DB	ID	Description
1	177.5	6.6	1105	2	Т47582	hypothetical prote
2	167.5	6.2	767	2	S63182	hypothetical prote
3	156	5.8	542	2	T46464	hypothetical prote
4	149	5.5	695	2	T40168	hypothetical prote
5	148.5	5.5	1403	1	A47328	natural killer cel
6	146.5	5.4	669	2	T28754	hypothetical prote
7	143	5.3	1577	2	T19722	hypothetical prote
8	143	5.3	3498	2	T22330	hypothetical prote
9	138.5	5.1	368	2	G88636	protein W09G12.7 [
10	135.5	5.0	643	2	A96636	unknown protein, 7
11	134.5	5.0	699	2	I38073	nucleolar phosphop
12	134.5	5.0	896	2	D96556	hypothetical prote
13	133.5	5.0	1672	2	T46237	hypothetical prote

14	133	4.9	705	2	D88536	acidic protein - C
15	133	4.9	705	2	S27786	acidic protein - C
16	133	4.9	943	2	A42681	centromere protein
17	131.5	4.9	425	2	S55147	hypothetical prote
18	130	4.8	608	2	T02299	hypothetical prote
19	130	4.8	679	2	S48437	hypothetical prote
20	129.5	4.8	2526	2	T20531	hypothetical prote
21	129.5	4.8	2722	2	T20532	hypothetical prote
22	129.5	4.8	2738	2	E88320	protein F07A11.6 [
23	128.5	4.8	543	2	T27190	hypothetical prote
24	128.5	4.8	552	2	T27191	hypothetical prote
25	128.5	4.8	954	2	E86174	protein F19P19.26
26	127.5	4.7	493	2	т02376	hypothetical prote
27	127.5	4.7	539	2	T15256	hypothetical prote
28	127	4.7	763	2	T08929	hypothetical prote
29	127	4.7	786	2	Т33856	hypothetical prote
30	127	4.7	845	2	A45669	neurofilament trip
31	127	4.7	963	2	T04002	hypothetical prote
32	126.5	4.7	390	2	T34137	hypothetical prote
33	126	4.7	598	2	B40713	cylicin I - human
34	126	4.7	1032	2	A57514	RNA helicase HEL11
35	125	4.6	1274	2	A89959	hypothetical prote
36	124.5	4.6	817	2	S53919	hypothetical prote
37	124	4.6	775	2	T21259	hypothetical prote
38	124	4.6	1166	2	н86341	hypothetical prote
39	123.5	4.6	849	2	E86306	Similar to tufteli
40	123	4.6	529	2	Т50609	hypothetical prote
41	122	4.5	581	2	T22455	hypothetical prote
42	122	4.5	611	2	T22456	hypothetical prote
43	122	4.5	971	2	T24866	hypothetical prote
44	122	4.5	1230	2	T22458	hypothetical prote
45	121	4.5	4910	2	S64942	probable membrane

#### ALIGNMENTS

```
RESULT 1
T47582
hypothetical protein F24B22.190 - Arabidopsis thaliana
C; Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 20-Apr-2000 #sequence revision 20-Apr-2000 #text change 09-Jul-2004
C; Accession: T47582
R; Bloecker, H.; Mewes, H.W.; Lemcke, K.; Mayer, K.F.X.; Quetier, F.; Salanoubat,
Μ.
submitted to the Protein Sequence Database, January 2000
A; Reference number: Z23016
A; Accession: T47582
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-1105 <BLO>
A; Cross-references: UNIPROT: Q9M383; UNIPARC: UPI00000A410D; EMBL: AL132957
A; Experimental source: cultivar Columbia; BAC clone F24B22
C; Genetics:
A; Map position: 3
A;Introns: 35/3; 56/2; 294/3; 318/3; 349/3; 376/2; 426/3; 455/1; 485/3; 508/3;
568/3; 633/1; 662/3; 681/3; 710/2; 981/1; 1043/3
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A; Note: F24B22.190

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# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame plus p2n model Run on: December 11, 2005, 17:38:44; Search time 5807 Seconds (without alignments) 4060.735 Million cell updates/sec Title: US-09-771-312-2 Perfect score: 2694 Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504 Scoring table: BLOSUM62 Xgapop 10.0 , Xgapext 0.5 Ygapop 10.0 , Ygapext 0.5 Fgapop 6.0 , Fgapext 7.0 Delop 6.0 , Delext 7.0 41078325 segs, 23393541228 residues Searched: Total number of hits satisfying chosen parameters: 82156650 Minimum DB seq length: 0 Maximum DB seq length: 2000000000 Post-processing: Minimum Match 0% Maximum Match 100% Listing first 45 summaries Command line parameters: -MODEL=frame+ p2n.model -DEV=xlh Q=/cgn2 1/USPTO spool/US09771312/runat 01122005 145312 15071/app query.fasta 1.6 -DB=EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000 -USER=US09771312 @CGN 1 1 5315 @runat 01122005 145312 15071 -NCPU=6 -ICPU=3 -NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG -DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7 Database : EST:\* 1: gb est1:\* 2: gb est2:\* 3: gb est3:\* 4: gb htc:\* 5: gb est4:\* 6: gb\_est5:\* 7: qb est6:\* 8: qb est7:\*

> 9: gb\_gss1:\* 10: gb\_gss2:\*

## 11: gb\_gss3:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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#### ALIGNMENTS

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            1
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            Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
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            Hubisz, M.J., Fledel-Alon, A., Tanenbaum, D.M., Civello, D.,
            White, T.J., Sninsky, J.J., Adams, M.D. and Cargill, M.
            A Scan for Positively Selected Genes in the Genomes of Humans and
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            2 (bases 1 to 1587)
            Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
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  JOURNAL
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REFERENCE
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           Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
           Hubisz, M.J., Fledel-Alon, A., Tanenbaum, D.M., Civello, D.,
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           A Scan for Positively Selected Genes in the Genomes of Humans and
 TITLE
           Chimpanzees
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           (er) PLoS Biol. 3 (6), E170 (2005)
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           White, T.J., Sninsky, J.J., Adams, M.D. and Cargill, M.
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 JOURNAL
           Submitted (05-MAY-2005) Celera Genomics, 45 West Gude Drive,
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           This sequence was made by sequencing genomic exons and ordering
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Qy	421	LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro	440
Db	1333	AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA	1392
Qу	441	<pre>IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer                                     </pre>	460
Db	1393	ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA	1452
Qy	461	GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys	480
Db	1453	GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG	1512
Qу	481	GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla	500
Db	1513	GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTGCTACCCCCAATGCA	1572
Qу	501	GlyLysSerAla 504	
Db	1573	GGAAAATCCGCC 1584	

#### us-09-771-312-2.rup

#### GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

December 4, 2005, 10:07:25; Search time 230 Seconds Run on:

(without alignments)
1546.027 Million cell updates/sec

Title: US-09-771-312-2

Perfect score: 2694

1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504 Sequence:

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

2166443 seqs, 705528306 residues Searched:

Total number of hits satisfying chosen parameters: 2166443

Minimum DB seq length: 0 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: UniProt\_05.80:\*

1: uniprot\_sprot:\* 2: uniprot\_trembl:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### **SUMMARIES**

Result No.	Score	% Query Match	Length	DB	ID	Description
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	2694 2694 2310 2009.5 1813 1587.5 1538 1513.5 1283.5 1008 939.5 627.5 593.5 545.5 438 430 320.5 197 177.5 177.5	100.0 100.0 85.7 74.6 67.3 58.9 57.1 567.6 37.4 34.9 23.3 22.2 16.3 11.9 7.3 6.6 6.3	528 528 527 504 376 414 375 410 561 216 408 482 453 467 107 221 928 1007 1105 812		GPTC2_HUMAN Q5VYK7_HUMAN GPTC2_MOUSE Q5F3Y2_CHICK Q5VYK8_HUMAN Q4V7S5_XENLA Q9D3E7_MOUSE Q6AY15_RAT Q4RRB2_TETNG Q6PIX0_HUMAN Q5RJ37_BRARE CN118_MOUSE Q9H3M3_HUMAN CN118_HUMAN Q4RLV5_TETNG Q9CSX3_MOUSE Q9ULA8_HUMAN Q6H4V9_ORYSA Q8VYR8_ARATH Q9M383_ARATH Q6C233_YARLI	Q9nw75 homo sapien Q5vyk7 homo sapien Q7tqc7 mus musculu Q5f3y2 gallus gall Q5vyk8 homo sapien Q4v7s5 xenopus lae Q9d3e7 mus musculu Q6ay15 rattus norv Q4rrb2 tetraodon n Q6pix0 homo sapien Q5rj37 brachydanio Q6pe65 mus musculu Q9h3m3 homo sapien Q9nwq4 homo sapien Q4rlv5 tetraodon n Q9csx3 mus musculu Q9ula8 homo sapien Q4rlv5 tetraodon n Q9csx3 mus musculu Q9ula8 homo sapien Q6h4v9 oryza sativ Q8vyr8 arabidopsis Q9m383 arabidopsis Q6c233 yarrowia li

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us-09-771-312-2.rup
                             742
         169
                   6.3
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166.5
23
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6.2
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Q55iv9 cryptococcu
Q9sf87 arabidopsis
24
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26
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732
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27
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28
29
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                                                                             059he6 homo sapien
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       162.5
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                                        Q99KV9_MOUSE
Q91YE7_MOUSE
Q6DDU9_XENLA
32
33
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Q6ddu9 xenopus lae
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157.5
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156
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Q9NTB1_HUMAN
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                                                                             Q99kg3 mus musculu
Q9ntb1 homo sapien
Q59ug4 candida alb
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41
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42
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                   5.8
                             853
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#### **ALIGNMENTS**

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05-JUL-2004 (Rel. 44, Created)
05-JUL-2004 (Rel. 44, Last sequence update)
10-MAY-2005 (Rel. 47, Last annotation update)
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DE
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GΝ
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OC.
OC.
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ox
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RN
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RC
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RA
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us-09-771-312-2.rup
         Nakagawa S., Senoh A., Mizoguchi H., Goto Y., Shimizu F., Wakebe H.,
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RA
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         cDNAs.":
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         Nat. Genet. 36:40-45(2004).
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         TISSUE=Lung, and Uterus;
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RA
RΑ
RA
        Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
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RA
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RA
         Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA
        Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human
and mouse cDNA sequences.";
RA
RA
RA
RT
RT
         Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RL
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CC
CC
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                    IsoId=Q9NW75-1; Sequence=Displayed;
CC
CC
                Name=2;
                    IsoId=Q9NW75-2; Sequence=VSP_010527, VSP_010528;
CC
CC
                    Note=No experimental confirmation available:
CC
         -!- SIMILARITY: Contains 1 G-patch domain.
CC
         This Swiss-Prot entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC
CC
         the European Bioinformatics Institute. There are no restrictions on its use as long as its content is in no way modified and this statement is not
CC
CC
CC
         removed.
CC
         EMBL; AK001114; BAA91509.1; -; mRNA. EMBL; BC042193; AAH42193.1; -; mRNA. EMBL; BC063474; AAH63474.1; -; mRNA.
DR
DR
DR
         Ensembl; ENSG00000092978; Homo sapiens.
DR
         HGNC; HGNC:25499; GPATC2.
DR
         InterPro; IPR000467; G_patch.
DR
         Pfam; PF01585; G-patch; 1.
SMART; SM00443; G_patch; 1.
DR
DR
DR
         PROSITE; PS50174; G_PATCH; 1.
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us-09-771-312-2.rup
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KW
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FT
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                      376
                               /FTId=VSP_010527.
Missing (in isoform 2).
FT
    VARSPLIC
               377
                      528
FT
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FT
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D -> N (in Ref. 2; AAH42193).
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    CONFLICT
               225
                      225
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                                472143144700DC26 CRC64;
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SQ
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100.0%; Pred. No. 1.5e-153;
tive 0; Mismatches 0;
                        100.0%;
 Query Match
                                                  Length 528;
 Best Local Similarity
 Matches 504: Conservative
                                                  Indels
                                                            0:
                                                               Gaps
                                                                       0:
QУ
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             Db
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Db
         121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qy
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             Db
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01-FEB-2005 (TrEMBLrel. 29, Last annotation update)
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Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae;
OC
OC
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OX
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    Griffiths C.;
RA
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EMBL; AC096641; CAH70664.1; JOINED; Genomic_DNA.
DR
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GO; GO:0003676; F:nucleic acid binding; IEA.
DR
DR
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SO
    SEQUENCE
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                                                                Gaps
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             Db
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Qу
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    GPTC2_MOUSE
                                PRT;
                                       527 AA.
    Q7TQC7; Q8BNJ9; Q8BPM1; Q8CDH9;
05-JUL-2004 (Rel. 44, Created)
05-JUL-2004 (Rel. 44, Last sequence update)
13-SEP-2005 (Rel. 48, Last annotation update)
G patch domain containing protein 2.
\mathsf{AC}
DT
DT
DT
DE
GN
    Name=Gpatc2;
os
    Mus musculus (Mouse).
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OC.
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OC.
oc
              Muroidea; Muridae; Murinae; Mus.
0X
              NCBI_TaxID=10090;
RN
              NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORMS 2 AND 3).
RP
RC
              STRAIN=C57BL/6J; TISSUE=Eye, and Testis;
              MEDLINE=22354683; PubMed=12466851; DOI=10.1038/nature01266;
RX
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RA
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RA
RA
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RA
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Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
Sandelin A. Schneider C. Semple C.A. Setou M. Shimada K.
RA
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ŘΑ
RA
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              Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A.,
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              Birney E., Hayashizaki Y.;
              "Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs.";
RT
RT
              Nature 420:563-573(2002).
RL
RN
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RP
              NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORM 1).
              STRAIN=C57BL/6; TISSUE=Brain; MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
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              Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
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Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
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              Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.
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             Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
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              Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A.,
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              Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
             Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
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Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human
RA
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              and mouse cDNA sequences.
              Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RL
              -!- ALTERNATIVE PRODUCTS:
\mathsf{CC}
                          Event=Alternative splicing; Named isoforms=3;
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CC
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CC
CC
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     This Swiss-Prot entry is copyright. It is produced through a collaboration
     between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use as long as its content is in no way modified and this statement is not
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CC
CC
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DR
     EMBL; AK053781; BAC35520.1; -; mRNA.

EMBL; AK083471; BAC38928.1; -; mRNA.

EMBL; BC054810; AAH54810.1; -; mRNA.

Ensembl; ENSMUSG00000039210; Mus musculus.
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DR
     MGI; MGÍ:1915019; Gpatc2.
DR
     InterPro; IPRO00467; G_patch.
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SMART; SM00443; G_patch; 1.
PROSITE; PS50174; G_PATCH; 1.
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S -> P (in Ref. 2).
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SQ
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84.9%; Pred. No. 1.7e-130;
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  Best Local Similarity
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            1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
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Qy
               84 GHCLSEGSDSSLEEPSKOYREKHSNNKKORSDSDDQMLVAKRRPSSNLSSSVRGKRLLWH 143
Db
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Qy
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Db
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